

The future of hyperdiverse tropical ecosystems

Jos Barlow¹, Filipe França^{1, 2, 3}, Toby A. Gardner⁴, Christina C. Hicks¹, Gareth D. Lennox¹, Erika Berenguer^{1, 5}, Leandro Castello⁶, Evan P. Economo⁷, Joice Ferreira³, Benoit Guénard⁸, Cecília Gontijo Leal⁹, Victoria Isaac¹⁰, Alexander C. Lees¹¹, Catherine L. Parr^{12, 13, 14}, Shaun K. Wilson^{15, 16}, Paul J. Young¹, Nicholas A. J. Graham¹

1. Lancaster Environment Centre, Lancaster University, Lancaster, UK

2. Instituto Federal de Minas Gerais, Bambuí, Brazil

3. Embrapa Amazônia Oriental, Belém, Brazil

4. Stockholm Environment Institute, Stockholm, Sweden

5. Environmental Change Institute, University of Oxford, Oxford, UK

6. Department of Fish and Wildlife Conservation, Virginia Tech, Blacksburg, USA

7. Okinawa Institute of Science and Technology Graduate University, Onna, Japan

8. School of Biological Sciences, The University of Hong Kong, Hong Kong SAR, China

9. Museu Paraense Emílio Goeldi, Belém, Brazil

10. Universidade Federal do Pará, Belém, Brazil

11. School of Science and Environment, Manchester Metropolitan University, Manchester, UK

12. School of Environmental Sciences, University of Liverpool, Liverpool, UK

13. University of Pretoria, Pretoria, South Africa

14. University of Witwatersrand, Wits, South Africa

15. Marine Science Program, Department of Biodiversity, Conservation and Attractions, Kensington, Australia

16. Oceans Institute, University of Western Australia, Crawley, Australia

Preface: The tropics contain the overwhelming majority of Earth's biodiversity: their terrestrial, freshwater and marine ecosystems hold over three-quarters of all species, including almost all shallow-water corals and >90% of terrestrial birds. Yet, tropical ecosystems are subject to pervasive and interacting stressors, such as deforestation, overfishing and climatic change. They are also set within a socio-economic context that includes growing pressure from an increasingly globalised world, larger and more affluent tropical populations, and the continuation of weak governance and limited response capacity. Concerted local, national and international actions are urgently required to prevent a collapse of tropical biodiversity.

Introduction

The tropics hold a disproportionate amount of global biological diversity, and are key to meeting the international community's aims of socially-just sustainable development and effective biodiversity conservation¹. Yet, tropical ecosystems are undergoing rapid environmental, socio-economic and demographic change², often driven by forces from extra-tropical, developed countries. The scale of these changes is unprecedented, and decisions implemented in the coming decades will define the future diversity and sustainability of the tropics.

Guiding these decisions depends on understanding the diversity and vulnerability of the four major tropical ecosystems: the forests and mesic savannas that cover most of the terrestrial tropics, the extensive freshwater systems that receive half of the world's rainfall, and the shallow-water coral reefs distributed along 150,000 km of coastline (Fig. 1). Here, we quantify and review the global importance of tropical biodiversity, evaluate the vulnerability of tropical

ecosystems to proximate stressors, and assess whether global and regional socio-economic changes will exacerbate or ameliorate biodiversity loss. We then examine the effectiveness of conservation approaches, and highlight the scientific advances required to foster positive change and help overcome the challenges arrayed against a sustainable tropical future.

The global importance of tropical ecosystems

Over evolutionary time, the tropics have acted both as a source and refuge for most extra-tropical terrestrial and marine species^{3,4}; but just how diverse and irreplaceable are the tropics today? The increase in species richness from polar to tropical regions, known as the latitudinal diversity gradient, repeats across a wide range of taxa and biomes. As a result of this gradient, tropical latitudes, which cover just 40% of the Earth's surface, hold a startling proportion of the planet's species: our assessment reveals that almost all shallow-water zooxanthellae corals, 91% of terrestrial birds, and >75% of amphibians, terrestrial mammals, freshwater fish, ants, flowering plants and marine fish have ranges that intersect tropical latitudes (Fig. 2a). For birds, the importance of the tropics extends far beyond 23.5 degrees of latitude, with almost half of all Nearctic species migrating to the Neotropics⁵ and over 2 billion passerine and near-passerines crossing the Sahara each autumn⁶. Moreover, a disproportionate number of the world's species are endemic to the tropics. For example, there are 4.5 times more endemic amphibians in the tropics than in temperate regions (Fig 2a). Tropical zones are less important for marine mammals and birds, taxa that peak in diversity at mid-latitudes^{7,8}. Nonetheless, >55% of these species use the tropics (Fig. 2a).

Overall, 78% of species across the ten taxa we assessed occurred within tropical latitudes, but incomplete taxonomic inventories mean that this is almost certainly an underestimate⁹. Between 15,000-19,000 new species are described annually¹⁰, and the majority of recently described terrestrial vertebrates¹¹ or predicted discoveries of invertebrates¹² are from the tropics. Even terrestrial mammals are still being discovered at a rate of c. 25 species a year, with the highest numbers in the Neo- and Afrotropics¹³. Shortfalls in species descriptions for other taxa are often far greater. For example, only 70,000 of an estimated 830,000 multi-cellular plants and animals have been named on coral reefs¹⁴, and the c. 500 spider species described each year represent a tiny fraction of the estimated 150,000 undescribed tropical species¹⁵.

Tropical taxonomic shortfalls are further compounded by a suite of systematic sampling biases. These include undersampling compared with temperate regions¹⁶, the spatial aggregation of sampling effort around coastal areas¹⁷, roads, rivers, urban settlements and high-profile research stations¹⁸, biases in favour of dry-season sampling when many invertebrate taxa are least abundant¹⁹, and the paucity of samples from ecosystems that are harder to access, such as mesophotic and rariphotic reefs²⁰.

The biological diversity of the tropics is mirrored by many forms of societal diversity²¹. For example, tropical countries contain 40% of the world's population yet 85% of extant languages are spoken within them²². The tropics also provide incalculable benefits to humanity. They house most of the key centres of plant domestication²³ and have been a vital laboratory for the development of science itself – the disciplines of ecology, biogeography and evolutionary

biology are founded on evidence gleaned from tropical ecosystems. Tropical ecosystems also make vital contributions to globally-important ecosystem services: covering just 0.1% of the ocean surface, coral reefs provide fish resources for 275 million people that live within 30 km of them²⁴ and coastal protection for up to 197 million people²⁵; humid tropical forests cover <12% of the world's ice-free land surface but produce 33% of global net primary productivity and store 25% of the carbon in the terrestrial biosphere²⁶; while tropical savannas provide a further 30% of net primary productivity and 15% of carbon storage²⁷. Tropical ecosystems also help drive vital atmospheric teleconnections. For instance, 70% of the rainfall in the 3.2M km² *Rio de la Plata* catchment is estimated to come from evaporation in Amazonia²⁸.

Vulnerability of tropical biota and ecosystems

For all five vertebrate groups with comprehensive IUCN assessments and spatial occurrence data²⁹, globally threatened species are more dependent on the tropics than those classed as Least Concern (Fig. 2b). In addition, 85% of species extinctions from these vertebrate groups have been of species that use the tropics²⁹. Consequently, although extinctions of other groups are less well understood, we can assume that most of the estimated 130,000 modern invertebrate extinctions³⁰ will also have been of tropical species. Thus, not only are the tropics vastly more diverse than temperate regions, this diversity is at far greater risk from human impacts³¹. Moreover, given that the tropics have the highest proportion of Data Deficient species and the lowest level of biodiversity-threat assessment¹⁶, information shortfalls mean we are likely underestimating the vulnerability of the tropical biome. We assessed this vulnerability in more depth by examining the effect of local and global stressors, the interactions between them, and the resulting changes to tropical ecosystems.

Local stressors

The tropics are subject to some of the highest rates of land-use change and degradation. While the spatial coverage of temperate forests has increased since 1990, tropical deforestation rates exceed 5M ha/yr³². Additional impacts stem from the expansion of large infrastructure projects (e.g. dams) and the growing demand for agricultural commodities, biofuels, timber, fuelwood and other natural resources³³. These all result in severe biotic responses. Even with mitigation, dams present a near-impassable barrier for river fish³⁴, while deforestation replaces a species-rich pool of forest-specialists with a smaller pool of common open-area species³⁵. The influence of land-use change also extends far into remaining natural areas through isolation and edge effects³⁶, additional anthropogenic disturbances³⁷ and altered climatic conditions³⁸. Edge effects suppress the abundance of threatened vertebrates up to 200-400 m into tropical forests³⁶, leaving almost no core forest refugia in the Brazilian Atlantic Forest where >80% is within 500 m of an edge³⁹. Even low levels of landscape modification have significant effects on range-restricted species³⁷, and time lags mean that some of the most deleterious effects are observed decades after landscape modification⁴⁰.

Pollution presents a diverse set of threats to tropical ecosystems. Inputs of sediments and nutrients from land-use change are well-established drivers of biodiversity loss across freshwater⁴¹ and coastal systems, including coral reefs⁴². Pesticide use is increasing across the tropics, reflecting rapid intensification of farming practices⁴³ and high pest pressures on tropical

crops⁴⁴. Tropical Asian rivers are a major source of the 1.2-2.4 million tonnes of plastic that enters the world's oceans each year⁴⁵, with micro-plastics entering into coral diets⁴⁶ and larger debris increasing rates of coral disease⁴⁷. These examples of chronic pollution are exacerbated by extreme events, such as of the *Fundão* Dam collapse, which released c. 50M m³ of waste into a 600 km stretch of river in south-east Brazil, causing a 7,000 km² toxic plume in the Atlantic Ocean⁴⁸.

Overexploitation is also pervasive across the tropics. Fishing has reduced fish biomass by over 75% across a third of coral reefs⁴⁹ and is shrinking the mean body size of exploited freshwater taxa⁵⁰. Hunting contributed to the loss of charismatic mega-herbivores, extirpating African elephants, rhinos and large predators from most of their original ranges^{51,52}. The world's tropical forests are affected by extensive over-harvesting of wildlife³¹, with estimates of the annual harvests of highly-trafficked animals such as pangolins reaching into the millions of individuals⁵³. Moreover, the growth in non-food uses of wildlife means that even small-bodied songbirds are at risk of global extinction⁵⁴. Overexploitation also extends beyond fauna and is driving economically valuable tropical trees to extinction⁵⁵.

Invasive species have been the second most important extinction driver of vertebrates since 1500 CE⁵⁶. Within terrestrial ecosystems, invasive species have exerted the strongest influence on islands and coastal mainlands⁵⁷, having driven thousands of species extinctions and altered trophic structures⁵⁸. On continents, they currently have a greater impact on economically developed and extra-tropical regions, but tropical ecosystems are predicted to become increasingly vulnerable to invasion in the 21st century⁵⁹. Despite a deficit of research in the tropics⁶⁰, two prominent examples highlight the scope and magnitude of species invasions into terrestrial tropical ecosystems: there has been an 84% increase of alien species detections between 2003 and 2010 in Singapore⁶¹, while invasive African grasses could threaten up to 380,000 km² of Australia's savannas by promoting landscape flammability⁶². In aquatic ecosystems, invasive predatory fishes, such as the Indo-Pacific lionfish in Caribbean coral reefs⁶³ and the Nile perch in African lakes⁶⁴, have contributed to the loss of native species. Marine invasions are also facilitated by the mass transport of species in ship ballast water, resulting in widespread biotic homogenisation⁶⁵.

Global climatic change

While many of the "local" stressors described above are promoted by globalised drivers, climate change is truly global. Increases in atmospheric CO₂ concentrations to levels >400 ppm has important implications for tropical terrestrial and aquatic ecosystems. Ocean acidification from dissolved CO₂ is changing ocean chemistry to the extent that declining coral calcification has already been detected⁶⁶. Conditions for reef accretion and growth may be mostly absent throughout the tropics by 2100 under business-as-usual emission scenarios⁶⁷. Within savannas, elevated CO₂ levels favour the growth of woody plants over grasses, contributing to woody encroachment and the potential for a switch in biome state^{68,69}. CO₂ fertilisation may have also contributed to enhanced tree productivity and mortality rates observed in humid tropical forests⁷⁰.

Global warming does not proceed at the same rate across the planet. Although the greatest absolute temperature increases are occurring at higher latitudes, the tropics are already some of the hottest places on the planet and have the lowest inter-annual temperature variability^{71,72}. Consequently, they will be the first areas to experience significantly warmer climates than the present day⁷² and will endure climatic conditions without present-day equivalents⁷¹. In addition, some of the most important climate oscillations, including El Niño and the Indian Ocean Dipole, take place within, and have their greatest influence on, tropical regions. It is unclear if these oscillations will change in a warming world, but extremes of their phases have the potential to exacerbate or ameliorate the overall warming trend. One outcome of increasing temperatures is the poleward shifts of species ranges or movement to higher altitudes or deeper depths⁷³. For example, corals in southern Japan are extending northwards at c. 14 km/yr⁷⁴, and temperate macroalgal communities are being replaced with corals and other tropical species along large stretches of Australian coastline⁷⁵. Latitudinal shifts in terrestrial and freshwater tropical species distributions are less certain because of the many natural and anthropogenic barriers, and the low dispersal capacity of many tropical species⁷⁶. Furthermore, the responses of terrestrial species are defined by changes in rainfall as well as temperature⁷⁷.

If movement is not an option, tropical species must adapt or face extinction. Unfortunately, there is evidence that some species are either approaching their physiological limits or are unable to adapt to the rate of environmental change⁷⁸. Increasing ocean temperature extremes are driving mass bleaching events and mortality of reef-forming corals, with the time between bleaching events declining by 76-80% since the early 1980s⁷⁹. Higher temperatures also affect tropical vertebrates, causing, for example, an extreme female bias in the sex ratio of green turtles in the warmer regions of the Great Barrier Reef⁸⁰ and a reduction in the reproductive success of African wild dogs⁸¹. Altered rainfall is also critical. Droughts are drying up biologically diverse small streams⁸², while even modest changes in dry-season length increase tropical tree mortality⁷⁰ and modify tropical forest bird community structure⁸³.

Stressor interactions and indirect effects

Stressors affecting tropical species can interact in myriad ways⁸⁴. We demonstrate this by compiling data from six case studies within a co-tolerance framework that allows species responses to two dominant stressors to be examined⁸⁵. Only a small subset of species or genera (8-32%) showed no or positive responses when both stressors were combined (Fig. 3), and up to 55% fell within the “double jeopardy” quadrant, indicating a negative response to both stressors. While our summary does not quantify the magnitude of effects, it clearly demonstrates that stressors can act together to reduce the abundance or occupancy of tropical species. Moreover, these co-tolerance analyses simplify the reality facing tropical ecosystems because most are affected by more than two stressors at any given location and time⁸⁴.

Many changes to tropical ecosystems result from indirect consequences of single or multiple stressors. On coral reefs, nutrient inputs from land may increase susceptibility to coral bleaching, disease, and outbreaks of pests⁸⁶, while poleward reef expansion is supported by feedbacks from range-shifting tropical herbivorous fish⁷⁵. Overexploitation can result in surprising changes in tropical ecosystem properties through trophic cascades. For instance, the

extirpation of a single detritivore fish species in the Orinoco basin reduced downstream organic-carbon transport, increasing net primary productivity and respiration⁸⁷. On reefs, overfishing of keystone predators has repercussions for benthic structure⁸⁸, while removal of herbivores can limit coral recovery from mass-mortality events⁸⁹. In mesic savannas, changes to herbivore numbers alter ecosystem functions and structure via their interactions with wildfire regimes⁹⁰. Invasive species are also frequently linked to other stressors: the introduction of the Nile perch played a major role in the decline of endemic fish species in Lake Victoria, but its effects were likely catalyzed by a combination of other drivers including soil erosion, eutrophication and overfishing⁶⁴.

Ecosystems in transition

Interactions between multiple anthropogenic stressors are causing pervasive changes in the tropics, such that alternate states are emerging across all major tropical ecosystems (Box 1). Perhaps counter-intuitively, trees are encroaching on savannas while grasses are invading disturbed tropical forests – but in both cases, changes are from species-rich to species-poor systems^{68,91}.

These drastic ecosystem transitions are accompanied by widespread modification of species composition. For example, the relative abundance of coral species has been altered on reefs that maintain coral dominance⁹²; extirpation of native fish has followed species introductions in lakes⁶⁴; liana biomass has increased in otherwise undisturbed Neotropical forests⁹³; and patterns of plant regeneration in humid forests have been altered by the overharvesting of seed-dispersing vertebrates^{31,94}. Altered species composition is a cause for concern because it could signal the onset of more severe modification, especially if dominant species are vulnerable or if there are cascading implications for ecosystem functioning. The collapse of Jamaican coral reefs provides one of the starkest examples. First, chronic overfishing depleted herbivorous fish populations, leaving the system over-reliant on sea urchins for grazing algae. Then Hurricane Allen impacted the system in 1980, creating a substantial amount of dead substrate. Although corals began recovering after the hurricane, the subsequent mass mortality of sea urchins due to disease, combined with the already low abundance of herbivorous fish, led to a phase shift from coral to macroalgal dominance^{95,96}.

Socio-economic context and response capacity

The interacting proximate stressors causing tropical environmental change are underpinned by broader changes in socio-economic and political factors. We examined the trajectories of four types of underlying distal drivers, including demography (Fig. 4a-b), socio-political factors (Fig. 4c-d), markets (Fig. 4e-f) and technology (Fig. 4g-h)⁹⁷ to explore how tropical countries are changing relative to the rest of the world and to evaluate the relative influence of local and global drivers. We also examined how the capacity of tropical countries to reduce or cope with proximate stressors compares to non-tropical countries based on underlying governance (Fig. 4i-j) and research capacity (Fig. 4k-l).

The immense biodiversity of the tropics exists in the context of rapid demographic and economic growth (Fig. 4a-b). Human population is growing at a faster rate in the tropics than

elsewhere (Fig. 4a) and by 2050 half of the world's population will live in the tropics². These demographic changes are accompanied by steady GDP growth, linked, in part, to the rapid expansion of agricultural and extractive industries. However, tropical per capita GDP – an important measure of human well-being – remains far lower than the non-tropical average (Fig. 4b), and the rates of change suggest little closing of the inequality gap between global south and north⁹⁸. Although the relationship between development and natural resource conservation does not have to be negative^{99,100}, measures reflecting higher social performance are almost always associated with higher resource use¹⁰⁰. A larger and more affluent tropical population will increase demands for timber, water, food, energy, and land, all of which are strongly linked with environmental degradation.

These internal changes will be exacerbated by economic growth in non-tropical countries, and the continued displacement of environmental impacts to less-developed areas¹⁰¹. Indeed, despite high levels of tropical cultural diversity^{21,22}, external socio-political influences (Fig. 4c-d) suggest that tropical countries have become increasingly susceptible to globalisation. For example, the proportion of imported food crops (Fig. 4c) and foreign-land acquisitions are far higher in the tropics than elsewhere (Fig. 4d) and are associated with extensive road building¹⁰² and agricultural investment¹⁰³. These trends towards increasing tropical globalisation are reinforced by changes in market integration (Fig. 4e-f) and technological development (Fig. 4g-h). For example, agricultural exports (Fig. 4e) are steadily increasing, albeit from a far lower baseline than the rest of the world. Moreover, given comparatively low levels of adoption of technological developments, such as industrial fishing techniques (Fig. 4g) or fertilizers (Fig. 4h), there is enormous risk that the rate of natural resource extraction in many tropical countries will increase further, supplying both domestic and export markets^{104,105}. Taken together, these examples highlight the crucial role that external markets will play in determining the fate of tropical ecosystems.

Effective environmental governance (Fig. 4i-j) is a necessary condition for improved sustainability outcomes¹⁰⁶, particularly when domestic (Fig. 4a-d) and global (Fig. 4c-f) distal drivers are expected to exert increasing and unsustainable pressure on tropical ecosystems^{2,103}. However, the World Bank's national-level assessments of governance effectiveness from the tropics sit in stark contrast to measures from extra-tropical countries, with no sign of improvement (Fig. 4i). External support for environmental governance may help where local governance is weak (Fig. 4j). Yet, despite greater OECD (Organisation for Economic Cooperation and Development) environmental aid in the tropics than elsewhere (Fig. 4j), these investments are dwarfed by the value of domestic resource extraction (e.g. agricultural exports; Fig. 4e), the value of which is two orders of magnitude greater than overseas environmental aid. Furthermore, OECD environmental aid has been declining in recent years and seems unlikely to increase in the short term¹⁰⁷.

Low governance capacity in the tropics is further exacerbated by insufficient research and development investment (Fig. 4k) and low levels of scientific output (Fig. 4l). Research investment is critical for driving innovation and the development of evidenced-based solutions to environmental degradation¹⁰⁸. Despite some notable centres of excellence, the vast majority of biodiversity-related data and research is concentrated in wealthy, non-tropical countries¹⁷

and manuscripts submitted by authors from low-income countries are less than half as likely to be published as those from high-income countries¹⁰⁹. These trends highlight an alarming disconnect between the global scientific process and the people that are most capable of engaging with decision makers, who have the best understanding of local context and, arguably, have the strongest incentive to achieve positive impacts through their research.

Diverse solutions for diverse systems

Tropical ecosystems – and therefore at least 78% of the world’s biodiversity (Fig. 2a) – are at a critical juncture. Multiple interacting local and global stressors (Fig. 3) that are driving species extinctions and potentially irreversible ecosystem transitions^{92,110} (Box 1) are set within a changing socio-economic context (Fig. 4). This changing context is characterised by growing and more affluent populations, an increasingly globalised world, and weak governance and research capacity – all of which threatens to increase environmental degradation, conflict and inequality¹⁰³. Countering these threats requires major improvements in local and global governance capacity and a step-change in how environmental objectives are integrated into broader development goals¹¹¹. We review the opportunities and limitations presented by three well-established and non-mutually exclusive approaches to conservation, before highlighting priorities for research.

Conservation approaches

A fundamental element of tropical conservation relies on protected areas to limit demographic pressures and the impact of local stressors. These are supported by a wealth of scientific evidence outlining the pervasive impact of local stressors across tropical ecosystems^{37,49} (Fig. 3) combined with an eco-centric philosophy that emphasizes the intrinsic rights of nature¹¹². Yet, despite significant expansion of protected-area coverage in the marine and forested tropics¹¹³, the current network remains poorly designed, has very limited coverage of tropical freshwaters and grasslands, and is inadequately resourced¹¹⁴. Moreover, a strategy focused solely on protected areas will not foster environmental conservation outside of reserves¹¹⁵ and fails to engage with the distal drivers of biodiversity loss (Fig. 4) that can undermine the effectiveness of protected areas themselves¹¹⁶.

A second set of approaches for tropical conservation is based on the notion that people need to perceive the benefits of nature to justify conservation. These emphasize the need to pursue conservation objectives in human-dominated landscapes, the provision of ecosystem services, and the involvement of private-sector actors. In the tropics, they are epitomised by the growth in market-based conservation payment mechanisms, such as REDD+¹¹⁷, investments in the “blue economy”¹¹⁸ and a step change in the number of companies making sustainability commitments¹¹⁹. These approaches have strengthened the conservation toolkit, especially where strict regulatory approaches have failed. Encouraging examples range from the positive effects of commodity certification (e.g. palm oil¹²⁰) to payment for ecosystem service schemes (e.g. watershed protection¹²¹). However, such approaches also attract significant criticism with implementation often lagging commitments¹¹⁹, persistent concerns around the social legitimacy of compensation schemes¹²², and the misalignment of market-based mechanisms with local needs and perceptions of environmental values¹²³.

A third and more diverse set of approaches is based on recognition of the interdependencies between people and nature, the coevolution of ecological and socio-economic systems at local, regional and global scales¹²⁴, and perspectives about the co-existence of people and nature. This set of more “systems-based” approaches includes: (1) an appreciation of the importance of bottom-up, community-based conservation approaches in human-dominated land- and seascapes (e.g. small-scale fisheries¹²⁵ and community-managed forests¹²⁶); (2) recognition of the role of indigenous people as environmental stewards, and shifts towards an appreciation of more collective relationships with nature (e.g. the Ecuadorian constitution¹²⁷); (3) landscape- and ecosystem-wide approaches that attempt to bridge the role of actors working at different scales and in different sectors (e.g. jurisdictional approaches to curb deforestation¹²⁸); and (4) a more explicit accounting of multi-scale feedbacks, including the role of distant market actors and distal drivers¹²⁴. These broad, multi-layered “people and nature” approaches hold considerable appeal, but the inherent complexity of local contexts can make them challenging to conceptualize, implement and measure in joined-up and consistent ways¹²⁹.

Acting together and acting now

The three broad approaches to the conservation and governance of tropical ecosystems outlined above are often associated with alternative researcher and practitioner worldviews^{130,131}. But the inherent ecological diversity (Fig. 2a), vulnerability (Figs. 2b & 3) and socio-economic complexity (Fig. 4) of the tropics highlights the importance of pluralism¹³² and the need to adopt a variety of what are often complementary and synergistic approaches¹³¹. For all their limitations, protected areas are indispensable to limit the impact of local stressors, and it will be impossible to avoid further biodiversity loss unless they are strengthened and expanded¹³³. However, conservation strategies must also address the underlying drivers of environmental change (Fig. 4) and avoid exacerbating deeply rooted inequalities¹¹⁵. Practice is always messier than theory, and the adoption of more sustainable management systems is usually only possible with the support of a range of actors, as can be seen in the recent successes of some hybrid governance approaches, with government, the private sector, and civil society organizations all playing vital roles¹³⁴.

Another clear message is that conservation efforts need to operate at local, regional and global scales to be effective. Many distal drivers are disconnected from sites of impact in both space and time, and the engagement of external actors, including in distant markets and governance processes, is often essential to ensure that local efforts are effective. These include more strategic integration of environmental policy with development goals¹³⁵, the need for multinational environmental governance approaches, especially for aquatic systems⁸², and recognition of the importance of tackling demand for unsustainable products from downstream buyers and investors¹¹⁹. The capstone of such efforts lies in the urgent need to deliver on the Paris Agreement, without which climate change will undercut or even negate hard-won local conservation successes, whether in coral reefs⁹² or tropical forests¹¹⁰.

Finally, we need to act now to address the pressing environmental challenges facing the tropics. This means being adaptive, learning by doing and embracing innovation. The last decades have seen a boom in proposals, innovations, and insights about the governance and management of

tropical ecosystems, ranging from more technocentric proposals to facilitate the evolution of climate-tolerant corals¹³⁶; ecological engineering to recover lost trophic interactions by species re-introductions, ecological replacements and rewilding¹³⁷; to radical new legal frameworks such as France's "Loi de vigilance" (2017-399) that places an unprecedented due diligence obligation on major companies to assess social and environmental risks in their supply chains beyond French borders. While these innovations serve different purposes and are varyingly scalable, they illustrate the potential of solutions-based science and conservation. Of course, acting now does not mean ignoring the existing evidence base or making uninformed decisions. Rather, it is vital that researchers and decision makers are vigilant to opportunities and risks and are willing to learn lessons.

Keeping pace with the Anthropocene

All approaches to governing tropical ecosystems will be more effective if they have legitimate local support and are based on strong scientific evidence that ensures, for example, that protected areas are located where they are most needed, ecosystem services are accurately quantified, extractive activities such as fishing and logging are managed sustainably, and underlying drivers of environmental degradation are identified and understood. Whilst these challenges are common to all conservation and sustainability science, they are magnified in the tropics due to their unique diversity, high vulnerability and the low research capacity of most tropical countries. Here, we examine four areas where research effort can be more closely aligned with some of the priorities highlighted by this review.

Addressing key knowledge shortfalls

Our understanding of tropical biodiversity is limited by significant knowledge shortfalls in taxonomy and species distributions¹³⁸. Overcoming these shortfalls will require targeting resources towards the information "black holes" that cover large regions of the tropics¹⁸. At the ecosystem level, there is a need for increased study of structurally and functionally distinct systems, particularly tropical grassy biomes⁶⁸, dry forests¹³⁹ and low-order stream systems¹⁴⁰. Progress in these areas will likely be aided by significant advances in DNA sequencing and informatics, which have the potential to invigorate taxonomic discovery, and reaching across cultural divides to incorporate national, regional and local knowledge that often remains ignored because it is not in English¹⁴¹, included in standard databases¹⁴², or recognised by conventional science¹⁴³.

Understanding vulnerability

Our growing knowledge of the role of individual stressors, such as landscape configuration or overexploitation, needs to be complemented by research on the impact of multiple stressors⁸⁴, which could help predict and mitigate complex biotic responses when climate and local stressors act in concert (Fig. 3). Other harder-to-study but important phenomena include the role of time lags or extinction debts⁴⁰, trophic cascades³¹, or trajectories of ecosystem degradation and recovery in the face of unprecedented environmental change¹⁴⁴. Revealing these more complex forms of vulnerability will often demand longer-term and larger multi-scale sampling and monitoring programs. New approaches are also needed to overcome one of the

409 more intractable challenges of tropical ecology: we often know least about the rarest and most
410 vulnerable species or taxonomic groups.

411 *Understanding distal drivers*

412 Conservation does not occur in a vacuum, and localised interventions are likely to be much
413 more effective if they are guided by a closer understanding of underlying distal drivers of
414 biodiversity loss and environmental change – including identifying the actors behind such
415 drivers, helping to determine potential trigger points and identifying more effective policy
416 responses⁹⁷. Unpicking the role of distal drivers is essential to understand how distant
417 interactions between social and environmental systems shape local environmental outcomes¹⁴⁵.
418 Careful study has revealed many surprising interactions, such as links between the
419 intensification of commercial fishing and increased bushmeat exploitation in west Africa¹⁴⁶, the
420 role of warfare in driving African mammal declines¹⁴⁷, or the role of exchange rates in driving
421 deforestation¹⁴⁸. Achieving this deeper understanding requires greater integration of the natural
422 and social sciences, with interdisciplinarity included as a core element of tropical-conservation
423 research¹⁴⁹.

424 *From research to impact*

425 Achieving positive impacts from conservation research relies on building a stronger science-
426 society interface that challenges the oversimplified assumption of a linear flow from knowledge
427 to action¹⁵⁰. Engendering positive changes will require closer participation of practitioners in the
428 research process and investments in outreach activities and professional capacity building¹⁵⁰.
429 These will be supported by studying the knowledge exchange process itself, including the critical
430 role played by knowledge brokers and boundary organizations^{151–153}. Part of this process will
431 involve a focus on success stories, or “bright spots”, enabling the social, institutional, and
432 environmental conditions that create positive outcomes to be identified and replicated¹⁵². The
433 positive social and ecological outcomes from innovative restoration and rewilding programmes
434 in Costa Rica and Mozambique demonstrate the potential for positive action¹⁵⁴.

435 Local managers and scientists have a vital role to play in designing and implementing research
436 that can inform regionally-appropriate conservation actions¹⁵⁵ – at present, our knowledge of
437 hyperdiverse ecosystems is over reliant on inferences gleaned from distant research stations or
438 inappropriate temperate theoretical constructs^{18,156}. Research is also more likely to have an
439 impact if the spatial scale of studies is more closely matched to the administrative scale at which
440 resource decisions are taken¹⁵⁷. Sustaining research programmes and learning networks in study
441 landscapes can also help build the vital relationships between researchers, local knowledge
442 holders and decision makers¹⁵⁵.

443 Achieving these changes requires building on trends in the technological, disciplinary and
444 cultural dimensions of research practice. In the technological domain, opportunities for data
445 collection have been revolutionised by developments in remote sensing and drones¹⁵⁸, the
446 plummeting costs of DNA technologies¹⁵⁹, and the step changes in bioinformatics that have
447 allowed big data to be stored and retrieved in open-access platforms¹⁶⁰. In the disciplinary
448 domain, the last decade has seen a marked uptick in inter- and transdisciplinary research, with a

greater – though still insufficient – integration of natural and social sciences. This has resulted in an increasing openness of researchers towards methodological pluralism and mixed-method approaches¹⁵⁰ and growing recognition of the contribution that can be made by local people, citizen- and para-scientists in biodiversity research¹⁶¹. Changes in research culture include the greater internationalisation of ecological science and closer approximation with society¹⁵⁰, both of which can help foster a more fertile ground for knowledge exchange and capacity building. Notable advances include the development of multi-disciplinary and multinational learning networks¹⁶², exponential growth in author teams¹⁶³, and major syntheses such as the Intergovernmental Platform for Biodiversity and Ecosystem Services (IPBES).

Recent years have seen a new awakening of environmental consciousness and calls for decisive action, manifest, for example, in the Paris Agreement, the Sustainable Development Goals, and voluntary Zero Deforestation Commitments. Tropical and non-tropical scientists can inform these endeavours by developing a reliable knowledge base and innovative management interventions. Overcoming the remaining research challenges is far from trivial and will require a massive investment of resources to develop scientific infrastructure and capacity within tropical nations, as well as profound changes to ways of working and the relationship between the research process and society at large. But a failure to act decisively and to act now will greatly increase the risk of unprecedented and irrevocable biodiversity loss in the hyperdiverse tropics.

Acknowledgements

For providing data, we thank Birdlife, International Union for Conservation of Nature, Ocean Biogeographic Information System, Charlie Veron for zooxanthellate corals, the Large-Scale Biosphere-Atmosphere Program (LBA) and National Environment Research Council grant NE/P004512/1 for forest birds, John Fell Fund for savanna birds, Pew Marine Fellows Program of The Pew Charitable Trusts for freshwater fish, and Brazilian Council for Scientific and Technological Development (CNPq) (PELD 441659/2016-0). Individual funding was provided by National Environment Research Council NE/K016431/1; NE/P004512/1; NE/L000016/1; European Research Council 759457; NASA's Interdisciplinary Research in Earth Sciences program (NNX14AD29G); OIST and JSPS KAKENHI (JP17K15180); CNPq scientific productivity grant (307788/2017-2); CNPq *Programa de Capacitação Institucional* (300231/2016-4); and Royal Society (UF140691).

Author contributions

JB developed the review with input from NAJG, TAG, CH, ACL and JF. FF and GDL analysed the data, supported by JB, TAG, CH, EB, LC, EPE, BG, CGL, VI, ACL, CLP, SW, PJY and NAJG. JB, TAG, CH, NAJG, LC, ACL, CLP, FF and GDL wrote the manuscript with input from all authors.

Reprints and permissions information is available at www.nature.com/reprints.

The authors declare no competing interests.

Correspondence and requests for materials should be addressed to josbarlow@gmail.com

487 Supplementary Information is linked to the online version of the paper at
488 www.nature.com/nature

489 **Figure legends**

490 **Figure 1 | The tropical biosphere. a,** Tropical terrestrial and marine biomes. The tropical
491 terrestrial biome (green) was defined as all tropical mesic ecoregions¹⁶⁴. These ecoregions span
492 82% of the 50 million km² of land between 23.5° N and 23.5° S, but extend into the subtropics in
493 some areas. The tropical marine biome was defined by the 1988-2018 mean minimum monthly
494 18 °C sea-surface isotherm. This isotherm bounds the latitudinal extent of shallow-water coral-
495 forming ecoregions (blue)¹⁶⁵. **b,** The Intertropical Convergence Zone (ITCZ). The ITCZ was defined
496 by 1979-2017 mid-summer (January – turquoise colour gradient – and July – red colour
497 gradient) mean monthly total rainfall >20 cm (where both January and July had rainfall >20 cm,
498 we show that with the largest total). The ITCZ is a strong predictor of the distribution of tropical
499 ecoregions (a). Data sources are presented in Extended Data Table 1.

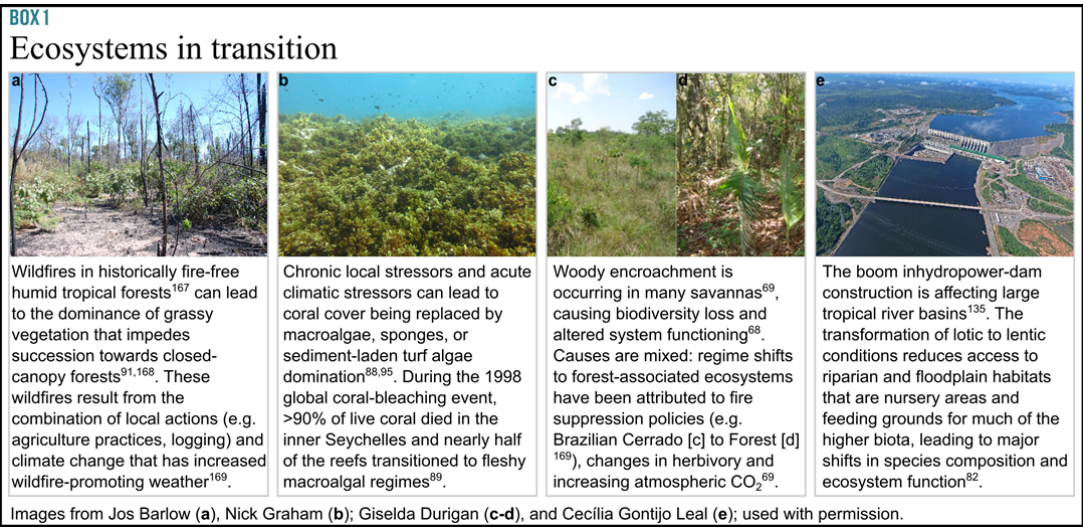
500 **Figure 2 | Tropical hyperdiversity. a,** The proportion of species found within tropical latitudes
501 for ten taxonomic groups. Bars are colour-coded to show the percentage of species ranges that
502 overlap the tropics. *n* gives the total number of species analysed in each group. Black boxes
503 around each bar show the proportion of all species that are endemic to the tropics. Only birds,
504 amphibians and mammals have been comprehensively sampled. Numbers at the end of the bars
505 give the precise percentage of species whose ranges overlap tropical latitudes, as shown in the
506 bars. **b,** The difference in the proportion of threatened (Critically Endangered, Endangered, and
507 Vulnerable) and non-threatened (Least Concern) species found exclusively within tropical
508 latitudes for the five comprehensively sampled groups. Data from: Birdlife International for
509 birds, the IUCN²⁹ for amphibians and mammals, the Ocean Biogeographic Information System
510 for marine fish, Charlie Veron for shallow-water zooxanthellate corals, Tedesco et al.¹⁶⁶ for
511 freshwater fish, and the Global Biodiversity Information Facility for angiosperms. Data sources
512 are presented in Extended Data Table 1.

513 **Figure 3 | Vulnerability of tropical biota to local and climatic stressors.** Species co-tolerance to
514 a local and climate-associated stressor⁸⁵. The x-axis shows responses to fishing for corals (a) and
515 reef (b) and freshwater fish (c); land-use change/deforestation for small-stemmed trees (2 ≤
516 DBH <10 cm; (d)) and forest birds (e); and fire suppression for savanna birds (f). The y-axis
517 represents longitudinal responses to climate-associated events: the 2015-16 and 1997-98 coral
518 bleaching events in the Seychelles for, respectively, corals (a) and reef fish (b); the 1997-98 El
519 Niño-induced drought for lower Amazonian freshwater fish (c); Amazonian fires during the
520 2015-16 El Niño for small-stemmed trees (d) and forest birds (e); and shrub encroachment
521 between 1998-2008 in South Africa for savanna birds (f). Species relative density is represented
522 from low (dark blue) to high (light green). The four quadrants represent the location of
523 “Survivor” species tolerant to both stressors (green), species only susceptible to local stressors
524 (yellow), species only vulnerable to climate-associated stressors (blue) and “double-jeopardy”
525 species susceptible to both stressors (red). Numbers show the percentage of species that fall
526 into the quadrant. *n* gives the total number of species – or genera for corals. Data sources are
527 presented in Extended Data Table 1.

528 **Figure 4 | Socio-economic drivers of biodiversity loss and societal response capacities.** Green
529 lines represent countries with >50% of their area within tropical latitudes; purple dashed-lines

represent all other countries; grey-shaded areas represent the proportion of the global total within tropical countries. **a**, Global population (1960-2016). **b**, Gross domestic product (GDP) per capita (2011 \$US based on purchasing power parity; 2000-2016). **c**, Foreign food crops (1961-2009). **d**, Cumulative overseas land ownership (2001-2017). **e**, Domestic and international airline passengers (1970-2016). **f**, Agricultural and forestry commodities export value (2001-2016). **g**, Bottom and pelagic trawler catch tonnages (1960-2014). **h**, Total fertilizer (nitrogen, potash, and phosphate) consumption relative to crop area (2002-2013). **i**, Government effectiveness index (2000-2016). **j**, Environmental protection aid (2000-2016). **k**, Public and private sector research and development expenditure (% GDP) (2000-2015). **l**, Scientific and technical journal articles per million people in the fields of physics, biology, chemistry, mathematics, clinical medicine, biomedical research, engineering and technology, and Earth and space sciences (2003-2016). Data sources are presented in Extended Data Table 1.

Box



Box text

Box 1. Tropical ecosystems in transition.

Forests (a): Wildfires in historically fire-free humid tropical forests¹⁶⁷ can lead to the dominance of grassy vegetation that impedes succession towards closed-canopy forests^{91,168}. These wildfires result from the combination of local actions (e.g. agricultural practices, logging) and climate change that has increased wildfire-promoting weather¹⁶⁹.

Corals (b): Chronic local stressors and acute climatic stressors can lead to coral cover being replaced by macroalgae, sponges, or sediment-laden turf algae^{89,95}. During the 1998 global coral-bleaching event, >90% of live coral died in the inner Seychelles and nearly half of the reefs transitioned to fleshy macroalgal regimes⁸⁹.

Savannas (c-d): Woody encroachment is occurring in many savannas⁶⁹, causing biodiversity loss and altered system functioning⁶⁸. Causes are mixed: regime shifts to forest-associated ecosystems have been attributed to fire suppression policies (e.g. Brazilian Cerrado [C] to Forest [D]¹⁷⁰), changes in herbivory and increasing atmospheric CO₂⁶⁹.

557 Freshwater (e): The boom in hydropower-dam construction is affecting large tropical river
558 basins¹³⁵. The transformation from lotic to lentic conditions reduces access to riparian and
559 floodplain habitats that are nursery areas and feeding grounds for much of the higher biota,
560 leading to major shifts in species composition and ecosystem function⁸².

561 References

- 562 1. SDG. Sustainable Development Goals. (2018). Available at:
563 <https://sustainabledevelopment.un.org/sdgs>. (Accessed: 20th January 2018)
- 564 2. Edelman, A. *et al. State of the Tropics: 2014 Report*. (2014).
- 565 3. Moreau, C. S. & Bell, C. D. Testing The Museum Versus Cradle Tropical Biological Diversity
566 Hypothesis: Phylogeny, Diversification, And Ancestral Biogeographic Range Evolution Of
567 The Ants. *Evolution (N. Y)*. **67**, 2240–2257 (2013).
- 568 4. Jablonski, D. *et al.* Out of the Tropics: Evolutionary Dynamics of the Latitudinal Diversity
569 Gradient. *Science (80-.)*. **314**, 102–106 (2006).
- 570 5. DeGraaf, R. M. & Rappole, J. H. *Neotropical Migratory Birds: natural history, distribution,*
571 *and population change*. (Cornell University Press, 1995).
- 572 6. Hahn, S., Bauer, S. & Liechti, F. The natural link between Europe and Africa - 2.1 billion
573 birds on migration. *Oikos* **118**, 624–626 (2009).
- 574 7. Tittensor, D. P. *et al.* Global patterns and predictors of marine biodiversity across taxa.
575 *Nature* **466**, 1098–1101 (2010).
- 576 8. Chown, S. L., Gaston, K. J. & Williams, P. H. Global patterns in species richness of pelagic
577 seabirds: the Procellariiformes. *Ecography (Cop.)*. **21**, 342–350 (1998).
- 578 9. Mora, C., Tittensor, D. P., Adl, S., Simpson, A. G. B. & Worm, B. How Many Species Are
579 There on Earth and in the Ocean? *PLoS Biol.* **9**, e1001127 (2011). **Develops a new**
580 **method to quantify the completeness of taxonomic inventories.**
- 581 10. Tancoigne, E. & Dubois, A. Taxonomy: no decline, but inertia. *Cladistics* **29**, 567–570
582 (2013).
- 583 11. Jenkins, C. N., Pimm, S. L. & Joppa, L. N. Global patterns of terrestrial vertebrate diversity
584 and conservation. *Proc. Natl. Acad. Sci.* **110**, E2602–E2610 (2013).
- 585 12. Guenard, B., Weiser, M. D. & Dunn, R. R. Global models of ant diversity suggest regions
586 where new discoveries are most likely are under disproportionate deforestation threat.
587 *Proc. Natl. Acad. Sci.* **109**, 7368–7373 (2012). **Identifies “hotspots of discovery” for ant**
588 **species.**
- 589 13. Burgin, C. J., Colella, J. P., Kahn, P. L. & Upham, N. S. How many species of mammals are
590 there? *J. Mammal.* **99**, 1–14 (2018). **Shows that the Neo- and Afrotropics contain the**
591 **highest number of newly recognised mammal species.**
- 592 14. Fisher, R. *et al.* Species richness on coral reefs and the pursuit of convergent global
593 estimates. *Curr. Biol.* **25**, 500–505 (2015).
- 594 15. Agnarsson, I., Coddington, J. A. & Kuntner, M. in *Spider research in the 21st century:*
595 *trends and perspectives* (ed. Penney, D.) 58–111 (Siri Scientific Press, 2013).
- 596 16. Collen, B., Ram, M., Zamin, T. & McRae, L. The tropical biodiversity data gap: Addressing
597 disparity in global monitoring. *Trop. Conserv. Sci.* **1**, 75–88 (2008).
- 598 17. Fisher, R. *et al.* Global mismatch between research effort and conservation needs of
599 tropical coral reefs. *Conserv. Lett.* **4**, 64–72 (2011).
- 600 18. Gardner, T. A. *et al.* Prospects for tropical forest biodiversity in a human-modified world.
601 *Ecol. Lett.* **12**, 561–582 (2009).
- 602 19. Barlow, J., Overal, W. L., Araujo, I. S., Gardner, T. A. & Peres, C. A. The value of primary,
603 secondary and plantation forests for fruit-feeding butterflies in the Brazilian Amazon. *J.*
604 *Appl. Ecol.* **44**, 1001–1012 (2007).
- 605 20. Baldwin, C. C., Tornabene, L. & Robertson, D. R. Below the Mesophotic. *Sci. Rep.* **8**, 4920
606 (2018).
- 607 21. Collard, I. F. & Foley, R. A. Latitudinal patterns and environmental determinants of recent
608 human cultural diversity: Do humans follow biogeographical rules? *Evol. Ecol. Res.* **4**,
609 371–383 (2002).
- 610 22. Simons, G. F. & Fennig, C. D. *Ethnologue: Languages of the World. Twenty-first edition.*
611 *Dallas, Texas: SIL International*. (2018). Available at: <http://www.ethnologue.com>.

- (Accessed: 20th February 2018)
23. Purugganan, M. D. & Fuller, D. Q. The nature of selection during plant domestication. *Nature* **457**, 843–848 (2009).
 24. Burke, L., Reyntar, K., Spalding, M. & Perry, A. *Reefs at risk revisited*. (2011).
 25. Ferrario, F. *et al.* The effectiveness of coral reefs for coastal hazard risk reduction and adaptation. *Nat. Commun.* **5**, 1–9 (2014).
 26. Bonan, G. B. Forests and climate change: Forcings, feedbacks, and the climate benefits of forests. *Science (80-.)*. **320**, 1444–1449 (2008).
 27. Grace, J. *et al.* Productivity and carbon fluxes of tropical savannas. *J. Biogeogr.* **33**, 387–400 (2006).
 28. Van Der Ent, R. J., Savenije, H. H. G. G., Schaeffli, B. & Steele-Dunne, S. C. Origin and fate of atmospheric moisture over continents. *Water Resour. Res.* **46**, 1–12 (2010). **Shows the importance of tropical forests for precipitation in extra-tropical regions.**
 29. IUCN. IUCN Spatial Data. (2018). Available at: <http://www.iucnredlist.org/technical-documents/spatial-data>.
 30. Régnier, C. *et al.* Mass extinction in poorly known taxa. *Proc. Natl. Acad. Sci.* **112**, 7761–7766 (2015).
 31. Dirzo, R. *et al.* Defaunation in the Anthropocene. *Science (80-.)*. **401**, 401–406 (2014).
 32. Keenan, R. J. *et al.* Dynamics of global forest area: Results from the FAO Global Forest Resources Assessment 2015. *For. Ecol. Manage.* **352**, 9–20 (2015).
 33. IUCN. *Threats Classification Scheme (Version 3.2)*. (2012).
 34. Pelicice, F. M., Pompeu, P. S. & Agostinho, A. A. Large reservoirs as ecological barriers to downstream movements of Neotropical migratory fish. *Fish Fish.* **16**, 697–715 (2015).
 35. Mendenhall, C. D., Shields-Estrada, A., Krishnaswami, A. J. & Daily, G. C. Quantifying and sustaining biodiversity in tropical agricultural landscapes. *Proc. Natl. Acad. Sci.* **113**, 14544–14551 (2016).
 36. Pfeifer, M. *et al.* Creation of forest edges has a global impact on forest vertebrates. *Nature* **551**, 187–191 (2017).
 37. Barlow, J. *et al.* Anthropogenic disturbance in tropical forests can double biodiversity loss from deforestation. *Nature* **535**, 144–147 (2016).
 38. Spracklen, D. V. & Garcia-Carreras, L. The impact of Amazonian deforestation on Amazon basin rainfall. *Geophys. Res. Lett.* **42**, 9546–9552 (2015).
 39. Haddad, N. M. *et al.* Habitat fragmentation and its lasting impact on Earth’s ecosystems. *Sci. Adv.* **1**, e1500052–e1500052 (2015).
 40. Gibson, L. *et al.* Near-complete extinction of native small mammal fauna 25 years after forest fragmentation. *Science (80-.)*. **341**, 1508–1510 (2013).
 41. Dudgeon, D. *et al.* Freshwater biodiversity: importance, threats, status and conservation challenges. *Biol. Rev.* **81**, 163 (2006).
 42. Altieri, A. H. *et al.* Tropical dead zones and mass mortalities on coral reefs. *Proc. Natl. Acad. Sci.* **114**, 3660–3665 (2017).
 43. Lewis, S. E., Silburn, D. M., Kookana, R. S. & Shaw, M. Pesticide Behavior, Fate, and Effects in the Tropics: An Overview of the Current State of Knowledge. *J. Agric. Food Chem.* **64**, 3917–3924 (2016).
 44. Bebber, D. P., Holmes, T. & Gurr, S. J. The global spread of crop pests and pathogens. *Glob. Ecol. Biogeogr.* **23**, 1398–1407 (2014).
 45. Lebreton, L. C. M. *et al.* River plastic emissions to the world’s oceans. *Nat. Commun.* **8**, 15611 (2017).
 46. Hall, N. M., Berry, K. L. E., Rintoul, L. & Hoogenboom, M. O. Microplastic ingestion by scleractinian corals. *Mar. Biol.* **162**, 725–732 (2015).
 47. Lamb, J. B. *et al.* Plastic waste associated with disease on coral reefs. *Science (80-.)*. **359**, 460–462 (2018).

- 663 48. Garcia, L. C., Ribeiro, D. B., de Oliveira Roque, F., Ochoa-Quintero, J. M. & Laurance, W. F.
664 Brazil's worst mining disaster: Corporations must be compelled to pay the actual
665 environmental costs. *Ecol. Appl.* **27**, 5–9 (2017).
- 666 49. MacNeil, M. A. *et al.* Recovery potential of the world's coral reef fishes. *Nature* **520**, 341–
667 344 (2015).
- 668 50. Castello, L. *et al.* The vulnerability of Amazon freshwater ecosystems. *Conserv. Lett.* **6**,
669 217–229 (2013).
- 670 51. Ripple, W. J. *et al.* Collapse of the world's largest herbivores. *Sci. Adv.* **1**, e1400103–
671 e1400103 (2015).
- 672 52. Ripple, W. J. *et al.* Status and Ecological Effects of the World's Largest Carnivores. *Science*
673 *(80-.)*. **343**, 1241484–1241484 (2014).
- 674 53. Ingram, D. J. *et al.* Assessing Africa-Wide Pangolin Exploitation by Scaling Local Data.
675 *Conserv. Lett.* **0**, 1–9 (2017).
- 676 54. Eaton, J. A. *et al.* Trade-driven extinctions and near-extinctions of avian taxa in Sundaic
677 Indonesia. *Forktail* **31**, 1–12 (2015).
- 678 55. Barrett, M. A., Brown, J. L., Morikawa, M. K., Labat, J.-N. & Yoder, A. D. CITES Designation
679 for Endangered Rosewood in Madagascar. *Science (80-.)*. **328**, 1109–1110 (2010).
- 680 56. Bellard, C., Cassey, P. & Blackburn, T. M. Alien species as a driver of recent extinctions.
681 *Biol. Lett.* **12**, 20150623 (2016).
- 682 57. Dawson, W. *et al.* Global hotspots and correlates of alien species richness across
683 taxonomic groups. *Nat. Ecol. Evol.* **1**, 186 (2017).
- 684 58. Simberloff, D. *et al.* Impacts of biological invasions: what's what and the way forward.
685 *Trends Ecol. Evol.* **28**, 58–66 (2013).
- 686 59. Early, R. *et al.* Global threats from invasive alien species in the twenty-first century and
687 national response capacities. *Nat. Commun.* **7**, 12485 (2016).
- 688 60. Pyšek, P. *et al.* Geographical and taxonomic biases in invasion ecology. *Trends Ecol. Evol.*
689 **23**, 237–244 (2008).
- 690 61. Nghiem, L. T. P. *et al.* Economic and Environmental Impacts of Harmful Non-Indigenous
691 Species in Southeast Asia. *PLoS One* **8**, e71255 (2013).
- 692 62. Setterfield, S. A., Rossiter-Rachor, N. A., Hutley, L. B., Douglas, M. M. & Williams, R. J.
693 Turning up the heat: the impacts of *Andropogon gayanus* (gamba grass) invasion on fire
694 behaviour in northern Australian savannas. *Divers. Distrib.* **16**, 854–861 (2010).
- 695 63. Albins, M. A. & Hixon, M. A. Worst case scenario: potential long-term effects of invasive
696 predatory lionfish (*Pterois volitans*) on Atlantic and Caribbean coral-reef communities.
697 *Environ. Biol. Fishes* **96**, 1151–1157 (2013).
- 698 64. Hecky, R. E., Mugidde, R., Ramlal, P. S., Talbot, M. R. & Kling, G. W. Multiple stressors
699 cause rapid ecosystem change in Lake Victoria. *Freshw. Biol.* **55**, 19–42 (2010).
- 700 65. Drake, J. M. & Lodge, D. M. Global hot spots of biological invasions: evaluating options
701 for ballast-water management. *Proc. R. Soc. B Biol. Sci.* **271**, 575–580 (2004).
- 702 66. Albright, R. *et al.* Reversal of ocean acidification enhances net coral reef calcification.
703 *Nature* **531**, 362–365 (2016).
- 704 67. Doney, S. C. *et al.* Climate Change Impacts on Marine Ecosystems. *Ann. Rev. Mar. Sci.* **4**,
705 11–37 (2012).
- 706 68. Parr, C. L., Lehmann, C. E. R., Bond, W. J., Hoffmann, W. A. & Andersen, A. N. Tropical
707 grassy biomes: Misunderstood, neglected, and under threat. *Trends Ecol. Evol.* **29**, 205–
708 213 (2014). **Highlights the importance of grassy tropical ecosystems.**
- 709 69. Stevens, N., Lehmann, C. E. R., Murphy, B. P. & Durigan, G. Savanna woody
710 encroachment is widespread across three continents. *Glob. Chang. Biol.* **23**, 235–244
711 (2017).
- 712 70. McDowell, N. *et al.* Drivers and mechanisms of tree mortality in moist tropical forests.
713 *New Phytol.* (2018). doi:10.1111/nph.15027

- 714 71. Williams, J. W., Jackson, S. T. & Kutzbach, J. E. Projected distributions of novel and
715 disappearing climates by 2100 AD. *Proc. Natl. Acad. Sci.* **104**, 5738–5742 (2007).
- 716 72. Mahlstein, I., Knutti, R., Solomon, S. & Portmann, R. W. Early onset of significant local
717 warming in low latitude countries. *Environ. Res. Lett.* **6**, 34009 (2011).
- 718 73. Pecl, G. T., Araújo, M. B., Bell, J. D., Blanchard, J. & ... Biodiversity redistribution under
719 climate change: Impacts on ecosystems and human well-being. *Science (80-.)*. **355**,
720 eaai9214 (2017).
- 721 74. Yamano, H., Sugihara, K. & Nomura, K. Rapid poleward range expansion of tropical reef
722 corals in response to rising sea surface temperatures. *Geophys. Res. Lett.* **38**, 1–6 (2011).
- 723 75. Wernberg, T. *et al.* Climate-driven regime shift of a temperate marine ecosystem. *Science*
724 *(80-.)*. **353**, 169–172 (2016).
- 725 76. Moore, R. P., Robinson, W. D., Lovette, I. J. & Robinson, T. R. Experimental evidence for
726 extreme dispersal limitation in tropical forest birds. *Ecol. Lett.* **11**, 960–968 (2008).
- 727 77. Vanderwal, J. *et al.* Focus on poleward shifts in species' distribution underestimates the
728 fingerprint of climate change. *Nat. Clim. Chang.* **3**, 239–243 (2013).
- 729 78. Hoffmann, A. A. & Sgrò, C. M. Climate change and evolutionary adaptation. *Nature* **470**,
730 479–485 (2011).
- 731 79. Hughes, T. P. *et al.* Spatial and temporal patterns of mass bleaching of corals in the
732 Anthropocene. *Science (80-.)*. **359**, 80–83 (2018).
- 733 80. Jensen, M. P. *et al.* Environmental Warming and Feminization of One of the Largest Sea
734 Turtle Populations in the World. *Curr. Biol.* **28**, 154–159.e4 (2018).
- 735 81. Woodroffe, R., Groom, R. & McNutt, J. W. Hot dogs: High ambient temperatures impact
736 reproductive success in a tropical carnivore. *J. Anim. Ecol.* **86**, 1329–1338 (2017).
- 737 82. Castello, L. & Macedo, M. N. Large-scale degradation of Amazonian freshwater
738 ecosystems. *Glob. Chang. Biol.* **22**, 990–1007 (2016).
- 739 83. Brawn, J. D., Benson, T. J., Stager, M., Sly, N. D. & Tarwater, C. E. Impacts of changing
740 rainfall regime on the demography of tropical birds. *Nat. Clim. Chang.* **7**, 133–136 (2017).
- 741 84. Côté, I. M., Darling, E. S. & Brown, C. J. Interactions among ecosystem stressors and their
742 importance in conservation. *Proc. R. Soc. B Biol. Sci.* **283**, 20152592 (2016). **Outlines the**
743 **importance of interactions between different stressors.**
- 744 85. Vinebrooke, R. D. *et al.* Impacts of multiple stressors on biodiversity and ecosystem
745 functioning: The role of species co-tolerance. *Oikos* **104**, 451–457 (2004).
- 746 86. Vega-Thurber, R. L. *et al.* Chronic nutrient enrichment increases prevalence and severity
747 of coral disease and bleaching. *Glob. Chang. Biol.* **20**, 544–554 (2014).
- 748 87. Taylor, B. W., Flecker, A. S. & Hall-Jr., R. O. Loss of a Harvested Fish Species Disrupts
749 Carbon Flow in a Diverse Tropical River. *Science (80-.)*. **313**, 833–836 (2006).
- 750 88. McClanahan, T. R. *et al.* Critical thresholds and tangible targets for ecosystem-based
751 management of coral reef fisheries. *Proc. Natl. Acad. Sci.* **108**, 17230–17233 (2011).
- 752 89. Graham, N. A. J., Jennings, S., MacNeil, M. A., Mouillot, D. & Wilson, S. K. Predicting
753 climate-driven regime shifts versus rebound potential in coral reefs. *Nature* **518**, 94–97
754 (2015).
- 755 90. Waldram, M. S., Bond, W. J. & Stock, W. D. Ecological Engineering by a Mega-Grazer:
756 White Rhino Impacts on a South African Savanna. *Ecosystems* **11**, 101–112 (2008).
- 757 91. Veldman, J. W. & Putz, F. E. Grass-dominated vegetation, not species-diverse natural
758 savanna, replaces degraded tropical forests on the southern edge of the Amazon Basin.
759 *Biol. Conserv.* **144**, 1419–1429 (2011).
- 760 92. Hughes, T. P. *et al.* Coral reefs in the Anthropocene. *Nature* **546**, 82–90 (2017).
- 761 93. Phillips, O. L. *et al.* Increasing dominance of large lianas in Amazonian forests. *Nature*
762 **418**, 770–774 (2002).
- 763 94. Harrison, R. D. *et al.* Consequences of defaunation for a tropical tree community. *Ecol.*
764 *Lett.* **16**, 687–694 (2013).

- 765 95. Hughes, T. P. Catastrophes, phase shifts and large-scale degradation of a Caribbean coral
766 reef. *Science (80-.)*. **265**, 1547–1551 (1994).
- 767 96. Mumby, P. J., Hastings, A. & Edwards, H. J. Thresholds and the resilience of Caribbean
768 coral reefs. *Nature* **450**, 98–101 (2007).
- 769 97. Hicks, C. C., Crowder, L. B., Graham, N. A. J., Kittinger, J. N. & Le Cornu, E. Social drivers
770 forewarn of marine regime shifts. *Front. Ecol. Environ.* **14**, 252–260 (2016).
- 771 98. Milanovic, B. Global Inequality: A New Approach for the Age of Globalization.
772 *Panoeconomicus* **63**, 493501 (2016).
- 773 99. Cinner, J. E. *et al.* Linking Social and Ecological Systems to Sustain Coral Reef Fisheries.
774 *Curr. Biol.* **19**, 206–212 (2009).
- 775 100. O'Neill, D. W., Fanning, A. L., Lamb, W. F. & Steinberger, J. K. A good life for all within
776 planetary boundaries. *Nat. Sustain.* **1**, 88–95 (2018). **Outlines the sustainability**
777 **challenges of current development trajectories.**
- 778 101. Stern, D. I., Common, M. S. & Barbier, E. B. Economic growth and environmental
779 degradation: The environmental Kuznets curve and sustainable development. *World Dev.*
780 **24**, 1151–1160 (1996).
- 781 102. Alamgir, M. *et al.* Economic, Socio-Political and Environmental Risks of Road
782 Development in the Tropics. *Curr. Biol.* **27**, R1130–R1140 (2017).
- 783 103. Deininger, K. & Byerlee, D. *Rising Global Interest in Farmland*. (The World Bank, 2011).
784 doi:10.1596/978-0-8213-8591-3
- 785 104. Laurance, Sayer, J. & Cassman, K. G. Agricultural expansion and its impacts on tropical
786 nature. *Trends Ecol. Evol.* **29**, 107–116 (2014).
- 787 105. Pauly, D. On Malthusian overfishing. *Naga, the ICLARM Quarterly* **13**, 3–4 (1990).
- 788 106. Rands, M. R. W. *et al.* Biodiversity Conservation: Challenges Beyond 2010. *Science (80-.)*.
789 **329**, 1298–1303 (2010). **Shows that effective environmental governance is a necessary**
790 **condition for improved sustainability outcomes.**
- 791 107. Blasiak, R. & Wabnitz, C. C. C. Aligning fisheries aid with international development
792 targets and goals. *Mar. Policy* **88**, 86–92 (2018).
- 793 108. Mora, C. *et al.* Management Effectiveness of the World's Marine Fisheries. *PLoS Biol.* **7**,
794 e1000131 (2009).
- 795 109. Mammides, C. *et al.* Increasing geographic diversity in the international conservation
796 literature: A stalled process? *Biol. Conserv.* **198**, 78–83 (2016). **Reveals that authors from**
797 **low-income countries are less than half as likely to be published as those from high-**
798 **income countries.**
- 799 110. Lovejoy, T. E. & Nobre, C. Amazon Tipping Point. *Sci. Adv.* **4**, eaat2340 (2018).
- 800 111. Nilsson, M. *Important interactions among the Sustainable Development Goals under*
801 *review at the High-Level Political Forum 2017*. (2017).
- 802 112. Kopnina, H., Washington, H., Gray, J. & Taylor, B. 'The "future of conservation" debate:
803 Defending ecocentrism and the Nature Needs Half movement'. *Biol. Conserv.* **217**, 140–
804 148 (2018).
- 805 113. ProtectedPlanet. World Database on Protected Areas (WDPA). (2018). Available at:
806 <https://www.protectedplanet.net/c/world-database-on-protected-areas>. (Accessed:
807 23rd February 2018)
- 808 114. Watson, J. E. M., Dudley, N., Segan, D. B. & Hockings, M. The performance and potential
809 of protected areas. *Nature* **515**, 67–73 (2014).
- 810 115. Büscher, B. *et al.* Half-Earth or Whole Earth? Radical ideas for conservation, and their
811 implications. *Oryx* **51**, 407–410 (2017).
- 812 116. Symes, W. S., Rao, M., Mascia, M. B. & Carrasco, L. R. Why do we lose protected areas?
813 Factors influencing protected area downgrading, downsizing and degazettement in the
814 tropics and subtropics. *Glob. Chang. Biol.* **22**, 656–665 (2016).
- 815 117. Agrawal, A., Nepstad, D. & Chhatre, A. Reducing Emissions from Deforestation and Forest

- Degradation. *Annu. Rev. Environ. Resour.* **36**, 373–396 (2011).
118. Winder, G. M. & Le Heron, R. Assembling a Blue Economy moment? Geographic engagement with globalizing biological-economic relations in multi-use marine environments. *Dialogues Hum. Geogr.* **7**, 3–26 (2017).
119. Lambin, E. F. *et al.* The role of supply-chain initiatives in reducing deforestation. *Nat. Clim. Chang.* **8**, 109–116 (2018). **Highlights the importance of tackling demand for unsustainable products from downstream buyers and investors.**
120. Carlson, K. M. *et al.* Effect of oil palm sustainability certification on deforestation and fire in Indonesia. *Proc. Natl. Acad. Sci.* **115**, 201704728 (2017).
121. Richards, R. C. *et al.* Governing a pioneer program on payment for watershed services: Stakeholder involvement, legal frameworks and early lessons from the Atlantic forest of Brazil. *Ecosyst. Serv.* **16**, 23–32 (2015).
122. Kosoy, N. & Corbera, E. Payments for ecosystem services as commodity fetishism. *Ecol. Econ.* **69**, 1228–1236 (2010).
123. Corbera, E. & Schroeder, H. REDD+ crossroads post Paris: Politics, lessons and interplays. *Forests* **8**, 1–11 (2017).
124. Liu, J. *et al.* Systems integration for global sustainability. *Science (80-.)*. **347**, 1258832–1258832 (2015).
125. Cinner, J. E. *et al.* Comanagement of coral reef social-ecological systems. *Proc. Natl. Acad. Sci.* **109**, 5219–5222 (2012).
126. Porter-Bolland, L. *et al.* Community managed forests and forest protected areas: An assessment of their conservation effectiveness across the tropics. *For. Ecol. Manage.* **268**, 6–17 (2012).
127. Gudynas, E. Buen Vivir: Today's tomorrow. *Development* **54**, 441–447 (2011).
128. Seymour, F. & Busch, J. *Why Forests? Why Now?: The Science, Economics, and Politics of Tropical Forests and Climate Change.* (Center for Global Development, 2016).
129. Mace, G. M. Whose conservation? *Science (80-.)*. **345**, 1558–1560 (2014).
130. Soulé, M. The 'new conservation'. *Keep. Wild Against Domest. Earth* **27**, 66–80 (2014).
131. Holmes, G., Sandbrook, C. & Fisher, J. A. Understanding conservationists' perspectives on the new-conservation debate. *Conserv. Biol.* **31**, 353–363 (2017).
132. Matulis, B. S. & Moyer, J. R. Beyond Inclusive Conservation: The Value of Pluralism, the Need for Agonism, and the Case for Social Instrumentalism. *Conserv. Lett.* **10**, 279–287 (2017). **Highlights the importance of pluralism in conservation approaches.**
133. Pouzols, F. M. *et al.* Global protected area expansion is compromised by projected land-use and parochialism. *Nature* **516**, 383–386 (2014).
134. Larsen, R. K. *et al.* Hybrid governance in agricultural commodity chains: Insights from implementation of 'No Deforestation, No Peat, No Exploitation' (NDPE) policies in the oil palm industry. *J. Clean. Prod.* **183**, 544–554 (2018).
135. Winemiller, K. O. *et al.* Balancing hydropower and biodiversity in the Amazon, Congo, and Mekong. *Science (80-.)*. **351**, 128–129 (2016).
136. van Oppen, M. J. H., Oliver, J. K., Putnam, H. M. & Gates, R. D. Building coral reef resilience through assisted evolution. *Proc. Natl. Acad. Sci.* **112**, 2307–2313 (2015).
137. Svenning, J.-C. *et al.* Science for a wilder Anthropocene: Synthesis and future directions for trophic rewilding research. *Proc. Natl. Acad. Sci.* **113**, 898–906 (2016).
138. Hortal, J. *et al.* Seven Shortfalls that Beset Large-Scale Knowledge of Biodiversity. *Annu. Rev. Ecol. Evol. Syst.* **46**, 523–549 (2015).
139. DRYFLOR *et al.* Plant diversity patterns in neotropical dry forests and their conservation implications. *Science (80-.)*. **353**, 1383–1387 (2016).
140. Leal, C. G. *et al.* Is environmental legislation conserving tropical stream faunas? A large-scale assessment of local, riparian and catchment-scale influences on Amazonian fish. *J. Appl. Ecol.* 1–15 (2017). doi:10.1111/1365-2664.13028

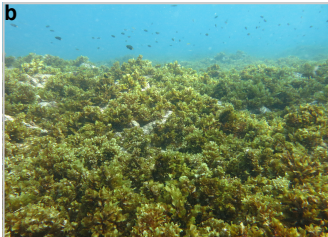
- 867 141. Pitman, N. C. A., Azádegui, M. D. C. L., Salas, K., Vigo, G. T. & Lutz, D. A. Written Accounts
868 of an Amazonian Landscape Over the Last 450 Years. *Conserv. Biol.* **21**, 253–262 (2007).
- 869 142. Feeley, K. Are We Filling the Data Void? An Assessment of the Amount and Extent of
870 Plant Collection Records and Census Data Available for Tropical South America. *PLoS One*
871 **10**, e0125629 (2015).
- 872 143. Sutherland, W. J., Gardner, T. A., Haider, L. J. & Dicks, L. V. How can local and traditional
873 knowledge be effectively incorporated into international assessments? *Oryx* **48**, 1–2
874 (2014).
- 875 144. Ghazoul, J. & Chazdon, R. Degradation and recovery in changing forest landscapes: A
876 multiscale conceptual framework. *Annu. Rev. Environ. Resour.* **42**, 161–188 (2017).
- 877 145. Liu, J. *et al.* Framing Sustainability in a Telecoupled World. *Ecol. Soc.* **18**, art26 (2013).
- 878 146. Brashares, J. S. Bushmeat hunting, wildlife declines, and fish supply in West Africa.
879 *Science (80-.)*. **306**, 1180–1183 (2004).
- 880 147. Daskin, J. H. & Pringle, R. M. Warfare and wildlife declines in Africa's protected areas.
881 *Nature* **553**, 328–332 (2018).
- 882 148. Richards, P. D., Myers, R. J., Swinton, S. M. & Walker, R. T. Exchange rates, soybean
883 supply response, and deforestation in South America. *Glob. Environ. Chang.* **22**, 454–462
884 (2012).
- 885 149. Hicks, C. C. *et al.* Engage key social concepts for sustainability. *Science (80-.)*. **352**, 38–40
886 (2016).
- 887 150. Fischer, J. *et al.* Advancing sustainability through mainstreaming a social–ecological
888 systems perspective. *Curr. Opin. Environ. Sustain.* **14**, 144–149 (2015). **Examines how the**
889 **concept of social-ecological systems can improve sustainability.**
- 890 151. Cvitanovic, C. *et al.* Improving knowledge exchange among scientists and decision-
891 makers to facilitate the adaptive governance of marine resources: A review of knowledge
892 and research needs. *Ocean Coast. Manag.* **112**, 25–35 (2015).
- 893 152. Cinner, J. E. *et al.* Bright spots among the world's coral reefs. *Nature* **535**, 416–419
894 (2016). **Shows how the "brightspots" approach can identify better than expected**
895 **environmental situations.**
- 896 153. Bennett, E. M. *et al.* Bright spots: seeds of a good Anthropocene. *Front. Ecol. Environ.* **14**,
897 441–448 (2016).
- 898 154. Pringle, R. M. Upgrading protected areas to conserve wild biodiversity. *Nature* **546**, 91–
899 99 (2017).
- 900 155. Balvanera, P. *et al.* Key features for more successful place-based sustainability research
901 on social-ecological systems: a Programme on Ecosystem Change and Society (PECS)
902 perspective. *Ecol. Soc.* **22**, 45 (2017).
- 903 156. Clarke, D. A., York, P. H., Rasheed, M. A. & Northfield, T. D. Does biodiversity–ecosystem
904 function literature neglect tropical ecosystems? *Trends Ecol. Evol.* **32**, 320–323 (2017).
- 905 157. Gardner, T. A. *et al.* A social and ecological assessment of tropical land uses at multiple
906 scales: the Sustainable Amazon Network. *Philos. Trans. R. Soc. B Biol. Sci.* **368**,
907 20120166–20120166 (2013).
- 908 158. Rose, R. A. *et al.* Ten ways remote sensing can contribute to conservation. *Conserv. Biol.*
909 **29**, 350–359 (2015).
- 910 159. Thomsen, P. F. & Willerslev, E. Environmental DNA - An emerging tool in conservation for
911 monitoring past and present biodiversity. *Biol. Conserv.* **183**, 4–18 (2015).
- 912 160. Gardner, T. A. *et al.* Transparency and sustainability in global commodity supply chains.
913 *World Dev.*
- 914 161. Basset, Y. *et al.* Conservation and biological monitoring of tropical forests: the role of
915 parataxonomists. *J. Appl. Ecol.* **41**, 163–174 (2004).
- 916 162. Barlow, J. *et al.* Using learning networks to understand complex systems: A case study of
917 biological, geophysical and social research in the Amazon. *Biol. Rev.* **86**, 457–474 (2011).

- 918 163. Barlow, J. *et al.* On the extinction of the single-authored paper: The causes and
919 consequences of increasingly collaborative applied ecological research. *J. Appl. Ecol.* **55**,
920 1–4 (2018).
- 921 164. Dinerstein, E. *et al.* An ecoregion-based approach to protecting half the terrestrial realm.
922 *Bioscience* **67**, 534–545 (2017).
- 923 165. Kleypas, J. A., McManus, J. W. & Meñez, L. A. B. Environmental Limits to Coral Reef
924 Development: Where Do We Draw the Line? *Am. Zool.* **39**, 146–159 (1999).
- 925 166. Tedesco, P. A. *et al.* A global database on freshwater fish species occurrence in drainage
926 basins. *Sci. Data* **4**, 170141 (2017).
- 927 167. Cochrane, M. A. Fire science for rainforests. *Nature* **421**, 913–919 (2003).
- 928 168. Flores, B. M., Fagoaga, R., Nelson, B. W. & Holmgren, M. Repeated fires trap Amazonian
929 blackwater floodplains in an open vegetation state. *J. Appl. Ecol.* **53**, 1597–1603 (2016).
- 930 169. Jolly, W. M. *et al.* Climate-induced variations in global wildfire danger from 1979 to 2013.
931 *Nat. Commun.* **6**, 1–11 (2015).
- 932 170. Durigan, G. & Ratter, J. A. The need for a consistent fire policy for Cerrado conservation.
933 *J. Appl. Ecol.* **53**, 11–15 (2016).
- 934

Ecosystems in transition



Wildfires in historically fire-free humid tropical forests¹⁶⁷ can lead to the dominance of grassy vegetation that impedes succession towards closed-canopy forests^{91,168}. These wildfires result from the combination of local actions (e.g. agriculture practices, logging) and climate change that has increased wildfire-promoting weather¹⁶⁹.



Chronic local stressors and acute climatic stressors can lead to coral cover being replaced by macroalgae, sponges, or sediment-laden turf algae^{88,95}. During the 1998 global coral-bleaching event, >90% of live coral died in the inner Seychelles and nearly half of the reefs transitioned to fleshy macroalgal regimes⁸⁹.



Woody encroachment is occurring in many savannas⁶⁹, causing biodiversity loss and altered system functioning⁶⁸. Causes are mixed: regime shifts to forest-associated ecosystems have been attributed to fire suppression policies (e.g. Brazilian Cerrado [c] to Forest [d]¹⁶⁹), changes in herbivory and increasing atmospheric CO₂⁶⁹.



The boom in hydropower-dam construction is affecting large tropical river basins¹³⁵. The transformation of lotic to lentic conditions reduces access to riparian and floodplain habitats that are nursery areas and feeding grounds for much of the higher biota, leading to major shifts in species composition and ecosystem function⁸².

aMinimum SST ($^{\circ}\text{C}$)

18-21

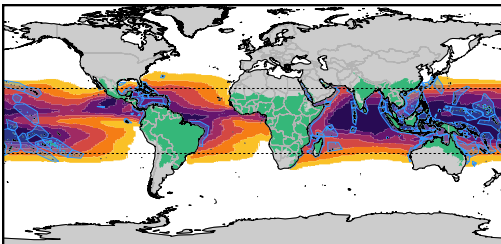
21-23

23-24

24-25

25-27

27-29

**b**

January

Rainfall (cm)

July

20

30

50

75

>150

20

30

50

75

>150

